

Phone: 888-417-5780 | Fax: 877-427-7290 | M-F, 8AM to 5PM EST |

# Please complete application in full, sign and date, then fax to: 877-427-7290

Or email to: ViatrisPAP@viatris.com

- The PAP Application must be complete to be reviewed for patient program eligibility. Please ensure all areas of the form are completed in full, including all signatures.
- To be considered for the Viatris Patient Assistance Program, all applicants must satisfy the following requirements and eligibility criteria:
  - o Applicants qualify for the program financial requirements.
  - Applicants must be a current United States resident (includes U.S Territories).
  - Applicants must be Uninsured.
  - The requested product must be prescribed by a licensed U.S. healthcare professional for a Food and Drug Administration (FDA) approved indication.
- Each applicant will be individually assessed for program eligibility based on the information provided within this application.
- Applicants will only be evaluated for eligibility upon receipt of a completed and signed Viatris Patient Assistance Program (PAP) Application.



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Patient information			
Name:	Date o	of Birth: / / SS	N:
Address*			
Home Phone: Cell P	hone: Patient Email	Address:	
Preferred Contact: Cell Phone Home Ph	one	ing Afternoon Evening	Gender:
Insurance: Uninsured Commercial	Government Other		
Insurance Name:	Insurance ID Number:	*No F	PO Boxes Accepted
Number of people in the household:	Gross Annual Household Income: (Including all Income, Wages, Social Security, Per	Gross Monthly Househnsion, Disability, Unemployment Benef	
Proof of Household Income will need to be submit Approved Verification Documents: 104 Advocate; or Certified Notarized States	0; 1040ez; W2; 4506-T; SSI Statement; Disabilit	ly Statement; Statement from Physic	cian, Nurse, or Patient
Prescriber Information			
Prescriber Name:		Prescriber NPI:	
Facility Name:		State License #:	
Facility Address:	City:	State:	ZIP:
Primary Office Contact:		Fax Number:	
Phone Number:	Office Contact Email:		
Prescriber Shipping Address (Only	complete if shipping address is diff	erent than address listed a	bove)
Prescriber Name:		Facility Name:	
Shipping Address:	City:	State:	ZIP:
Shipment Contact Name:			
Phone Number:	Contact Email:		



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Ohio Prescriber Mandatory Subsection (Select an option below, complete the related fields, then sign & date)!

MANDA	TORY SUBSECTION FOR ALL OHIO HCPs	
as a Terr TDDD lic For more www.pha	minal Distributor of Dangerous Drugs ("TDDD") o cense allows a business entity to receive, purchas e information on TDDD licensing requirements for	impany, may only provide prescription drugs to a prescriber whose practice is licensed in is exempt from such licensure under Ohio Revised Code ("ORC") § 4729.541. A see, and possess prescription drugs, including drug samples, for distribution to patients. It is prescribers, please visit the Ohio Board of Pharmacy website at fexemptions, please refer to section 4729.541 of the ORC. The above information is or should it be construed, as legal advice.
Please s	select and complete one of the following and s	sign below:
	The practice at which I work,	, located at the address I provided above, has an active TDDD license that
		rescription drug products at this location. The TDDD license number is
	which exp	ires on
-OR-		
	The practice at which I work,	, located at the address I provided above, is subject to one of the TDDD
	licensing exemptions in ORC § 4729.541.	<del></del>
prescript	•	above is complete and accurate and attest that I can receive and store the requested use I hold an unrestricted, active TDDD license or my practice is exempt from obtaining
Prescribe	er Signature:	Date:
	(Original signature -and	l- date required, stamped signatures not accepted)



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#### Product & Prescription Information (Select a Product & Complete Rx Details)

Rx Product Quantity Select	ion- Please indicate a selection below	by adding <b>quantity</b> to comple	te the Prescription
Arixtra® (fondaparinux sodium)	EMSAM® Transdermal System	Luxiq®	Proctofoam® HC
injection, solution	12 mg/24 hr Bx30	(betamethasonevalerate) Foam	(hydrocortisone acetate 1% and
2.5mg/0.5mL PFS 10PK	QTY	0.12% 100gm	pramoxine hydrochloride 1%)
QTY	Q11	QTY	HC 1% 10g
5mg/0.4mL PFS 10PK	6 mg/24 hr Bx30	0.12% 50gm	QTY
QTY	QTY	QTY	ROWASA® (mesalamine) Recta
7.5mg/0.6mL PFS 10PK	TDS 9 mg/24 hr Bx30		Suspension
QTY		Miacalcin® Injection	60mL Rectal Susp 7s
10mg/0.8mL PFS 10PK	QTY	200 IU/mL 2mL MDV	QTY
QTY	Erygel® (erythromycin) Topical Gel USP,	1pk	60mL Rectal Susp 28s
Cim duo® (lamivudine and	2%	QTY	QTY
tenofovir disoproxil fumarate)	2% 30gm	Muse® (alprostadil) urethral	- (POWA OA ® / Lucia )
tablet	QTY	250mcg Suppository 6s	sfROWASA® (mesalamine) Rectal Suspension
300/300mg T 30s		QTY 230116g Suppository 03	rectal caspension
QTY	2% 60gm		60mL Rectal Susp 7s
Cortifoam® (hydrocortisone	QTY	500mcg Suppository 6s	QTY
acetate 10%) rectal foam	Evoclin® (clindamycin phosphate) Foam,	QTY	60mL Rectal Susp 28s
10% 15g	1%	1000mcg Suppository	QTY
QTY	10/ 100gm	6s	Semglee® (Insulin Glargine)
Cystagon® (Cysteamine	1% 100gm	QTY	Injection
bitartrate) capsules	QIY	Olux® (clobetasol propionate)	1000 IU/10mL (1 Vial)
50mg C 500s	1% 50gm	Foam, 0.05%	QTY TOOO IO/ TOTTLE (1 VIAI)
QTY	QTY	0.05% 50gm	
150mg C 500s	<b>7</b> II 4 10 (5 II 4 )	QTY 0.03 % 30gm	300 IU/3mL (5 PF Pens)
QTY	Felbatol® (felbamate)		QTY
Denavir® (penciclovir) Cream	400mg T 100s	0.05% 100gm	Wixela Inhub® (fluticasone
1% 5gm	QTY	QTY	propionate and salmeterol
QTY	000 T 100	Olux-E® (clobetasol propionate)	inhalation pow der, USP)
Depen® (penicillamine tablets,	600mg T 100s	Foam, 0.05%	100mcg/50mcg 60/lnh
USP) Titratable Tablets	QTY	0.05% 50gm	QTY
250mg T 100s	600mg OS 8oz	QTY 0.03 % 30gm	250mcg/50mcg 60/lnh
QTY	QTY		QTY
Dipentum® (olsalazine sodium)		0.05% 100gm	500mcg/50mcg 60/lnh
capsule	600mg OS 32oz	QTY	QTY
250mg C 100s	QTY	Perforomist® (formoterol	
QTY	Gastrocrom® (cromolyn sodium, USP)	fumarate) Inhalation Solution	XULANE® (norelgestromin and ethinyl estradiol transdermal
Dym ista® azelastine	oral concentrate	,	system)
hydrochloride & fluticasone		20 mcg / 2 mL 30x1	TDS 0.15mg/0.035mg/Q
propionate) nasal spray		QTY	3s
137/50mcg Nasal Spray 23g	100mg 5mL Oral Concentrate 96s	20 mcg / 2 mL 60x1	QTY
QTY	QTY	QTY	V 1101
	Impeklo® (clobetasol propionate)	Pretomanid Tablets	Yupelri® (revefenacin) inhalation solution
Elestrin® (estradiol gel) 0.06%	Lotion MDP 0.05%		
0.06% Gel 35g Dual Pack		200mg T 26	175mcg / 3mL 30s
QTY	QTY	QTY	QTY



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Prescription Details- Please complete all relevant prescription details below
Patient Name:Patient DOB:
Prescriber Name:Prescriber NPI:
Day Supply:Refills:
Directions:
Prescriber Certification and Prescription Signature
I certify that the information provided in this Patient Assistance Program Application is complete and accurate to the best of my knowledge, that the Viatris product I have prescribed to the applicant within this application is based on my professional judgment of medical necessity for a Food and Drug Administration (FDA) approved indication, and that I will supervise the patient's medical treatment. I will notify Viatris PAP immediately if the Viatris product is no longer medically necessary for this patient's treatment. I certify that I have obtained from my patient all required written authorizations for the release of my patient's personal identification and insurance information to Viatris and their agents and representatives.
I understand that any information provided to Viatris and its agents and representatives is for the sole use of Viatris and their agents, service providers and representatives to verify my patient's insurance coverage status, to assess the patient's eligibility for participation in the Viatris Patient Assistance Program (collectively, "the Program"), and to otherwise administer the product and related services. I understand that application to the Program does no guarantee that assistance will be obtained.
I understand that Viatris may change or cancel this program at any time. I understand that if my patient's financial and/or insurance status changes, the patient may no longer be eligible for the Program, and I agree to immediately notify a Viatris PAP representative if I become aware of changes in the patient's financial and/or insurance status. I agree that Viatris PAP may contact me for additional information relating to this application either by fax, e mail and/or telephone. I understand that I am under no obligation to prescribe any Viatris product and that I have not received, nor will I receive, any benefit from Viatris or its agents or representatives for prescribing a Viatris product. I agree that I will not sell, submit claims or make any attempt to received reimbursement from any third party for any product provided by the Program.
Prescriber acknowledges that in connection with the application and enrollment process, United BioSource Corporation (UBC) performs eligibility screening using the Surescripts network. Surescripts requires that Prescriber agree to comply with all Surescripts' terms and conditions, including confidentiality commercial messaging, privacy and security, applicable laws, and use of data. All Surescripts disclaimers apply. A full list of terms and conditions is available at https://ubc.com/surescriptsterms/.
By signing this Patient Assistance Program Application, I authorize the release of medical and/or other patient information to agents and service providers of Viatris to use and disclose as necessary for verification of patient eligibility, and to furnish any information on this form to the insurer of the applicant for the purpose of verifying benefit eligibility. I understand that Program duration per eligibility period is 12 months, and the maximum number of refills per eligible patient is 11 for each unique enrollment.
Prescriber Certification & Prescription Signature: Date:
(original signature required)



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#### **Patient Authorization and Agreement Signature**

By signing this Authorization, I authorize each of my physicians, pharmacists, including any non-commercial pharmacy that receives my prescription ("my Prescribed Product"), and other healthcare providers (together "Healthcare Providers") and each of my health insurers, if any (together, "Insurers") to disclose my Protected Health Information, including but not limited to medical records, information related to my medical condition and treatment, my health insurance coverage, my name, address, telephone number, Social Security number, insurance plan and or group numbers (together, "Protected Health Information") to Viatris, its affiliated companies, vendors, agents, collaboration partners, and representatives (together, "Viatris") including providers of alternate sources of funding for prescription drug costs, and other service providers supporting the Viatris Patient Assistance Program (PAP) (collectively, the "Program") for the purposes described below.

Specifically, I authorize disclosure of my Protected Health Information in order to:

- I. Enroll me in, and contact me about the Program, including online support, financial assistance services, and co-pay assistance services, as applicable,
- II. Communicate with my Healthcare Providers and Insurers about benefits, coverage, and medical care, including compliance with Product treatments,
- III. Facilitate dispensing of my prescription by a non-commercial pharmacy,
- IV. Provide me with educational materials, information and services related to my treatment experience with my prescribed medication and my condition,
- V. Verify, investigate, and coordinate with my Insurers regarding my prescribed medication, and
- VI. Contact me as otherwise required or permitted by law.

Once my Protected Health Information has been disclosed to Viatris, I understand that federal privacy laws no longer protect the information. However, Viatris agrees to protect my Protected Health Information by using and disclosing it only for the purposes described in this Authorization or as permitted by law. I understand that I may refuse to sign this Authorization. My choice about whether to sign will not change the way my Healthcare Providers or Insurers treat me, but I will not have access to the Viatris Patient Assistance Program and the services provided by Viatris under the Program. If I refuse to sign the Authorization, or revoke my Authorization later, I understand that this means I will not be able to participate in or receive assistance from the Program.

I understand that my signed Authorization is valid for 5 years from the date of my signature, and that I may revoke this Authorization at any time in the future, except to the extent that actions have been taken in reliance on the Authorization. I understand that to revoke this Authorization I may mail a request to 5005 Greenbag Road Morgantown, WV 26508, fax to 877-427-7290, or by calling 888-417-5780. I understand that revoking this Authorization will end further uses and disclosure of my Protected Health Information by the parties identified above except to the extent those uses and disclosures have been made in reliance upon this Authorization as permitted by applicable law. I am entitled to receive a copy of this Authorization.

I understand that if I qualify and I am enrolled in the Program sponsored by Viatris, I will receive my Prescribed Product from Viatris only pursuant to a legally valid prescription from my health care provider. I understand that if I qualify and I am enrolled in the Program, Viatris will provide me my Prescribed Product free of charge for the duration of the enrollment period so long as I have a legally valid prescription for my Prescribed Product. I understand that I am not required to continue treatment with my Prescribed Product if I gain insurance coverage, or to receive treatment from any given provider. I understand and agree that I must notify Viatris PAP at 888-417-5780 immediately if my insurance status changes during the Program enrollment period. I understand and agree that neither I nor my Insurers, if applicable, will be charged for the supply of my Prescribed Product that I received from the Program, and that under NO circumstances may I claim reimbursement from my Insurers or any other third party for the Prescribed Product provided to me free of charge from the Program. I understand that Viatris reserves the right at any time without notice to modify or discontinue the Program and its criteria.

I understand that I am providing 'written instructions' to Viatris under the Fair Credit Reporting Act authorizing Experian on behalf of Viatris to obtain information from my credit profile or other information from Experian. I authorize Viatris and its service providers to obtain such information solely for the purpose of determining financial qualifications for the Program. I understand that I must affirmatively agree to the terms in this notice by signing below in order to proceed in the Program financial screening process.

My signature certifies that I have read and understand the above statements and agree to the outlined terms.

Patient Name (Print):	Patient Signature	: Date:	_
Patient Authorized Representati	ive		
	estions, any missing documentation	ring person about this application form. This includes discussing the status on and other issues related to my enrollment, or any other treatment-relate my time by calling: 888-417-5780	
Name of Authorized Representative:		Relationship to Patient:	_
Telephone Number:	Email:		_
By signing below, I, the patient, allow this re	epresentative to speak on my beha	alf on any matter regarding my enrollment with the Program.	
Patient Signature:		Date:	

